ALLERGY MEDICINE- diphenhydramine hcl tablet, film coated COSTCO WHOLESALE CORPORATION

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Kirkland 44-329

Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - runny nose
 - itchy, watery eyes
 - sneezing
 - itching of the nose or throat
- temporarily relieves these symptoms due to the common cold:
 - runny nose
 - sneezing

Warnings

Do not use

- to make a child sleepy
- with any other product containing diphenhydramine, even one used on skin

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- marked drowsiness may occur
- avoid alcoholic beverages

- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- do not take more than directed
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

	1 to 2
over	tablets
children 6 to under 12 years	1 tablet
children under 6 years	do not use

Other information

- each tablet contains: calcium 30 mg
- protect from moisture
- use by expiration date on package
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

corn starch, D&C red #27 aluminum lake, dibasic calcium phosphate dihydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

KIRKLAND

Signature

COMPARE TO BENADRYL® ALLERGY ULTRATAB® TABLETS

active ingredient*

NDC 63981-329-78

ITM. / ART. 536099

ALLERGY MEDICINE

Diphenhydramine HCl 25 mg

Antihistamine • Allergy Relief

For Temporary Relief of: Sneezing & Runny Nose Itchy, Watery Eyes Hay Fever Symptoms Itching of Nose or Throat

Actual Size

600 Minitabs

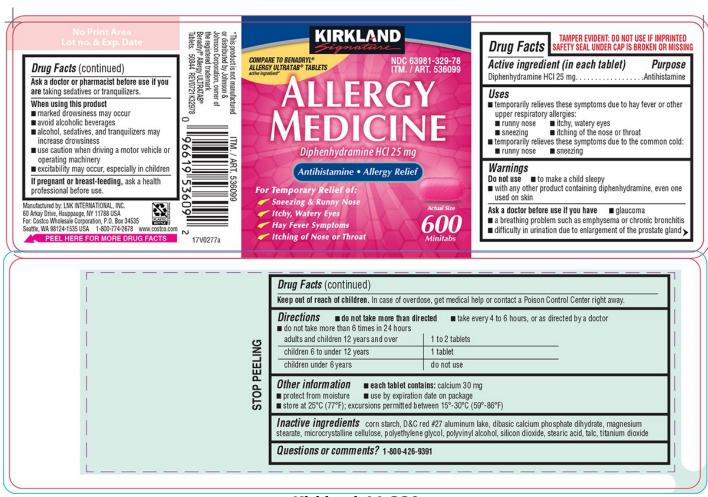
TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Benadryl® Allergy ULTRATAB® Tablets. 50844 REV0721K32978

Manufactured by: LNK INTERNATIONAL, INC. 60 Arkay Drive, Hauppauge, NY 11788 USA

For: Costco Wholesale Corporation, P.O. Box 34535

Seattle, WA 98124-1535 USA 1-800-774-2678 www.costco.com



Kirkland 44-329

ALLERGY MEDICINE

diphenhydramine hcl tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63981-329
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

STEARIC ACID (UNII: 4ELV7Z65AP)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics			
Color	pink	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	44;329
Contains			

	Packaging				
-	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:63981- 329-78	600 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/1990	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part341	03/02/1990			

Labeler - COSTCO WHOLESALE CORPORATION (103391843)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63981-329)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63981-329) , pack(63981-329)

Establishment			
Name	Address	ID/FEI	Business Operations
INV International Inc		022067004	manufacturo/62001 220)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(63981-329)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(63981-329)

Revised: 4/2023 COSTCO WHOLESALE CORPORATION